



Medicines & Healthcare products
Regulatory Agency



MHRA
10 South Colonnade
London
E14 4PU
United Kingdom
www.gov.uk/mhra

06th January 2023

Dear

FOI 22/1206

Thank you for your email, dated 16th December 2022, in which you requested:

- *Current approved: Risk Management Plan, RMP (eCTD Module 1.8.2 Risk Management System) for PLGB 00101/1223 Zolgensma 2 x 10¹³ vector genomes/mL solution for infusion*

We can confirm that the MHRA holds a copy of the current approved RMP (Version 1) and it has been attached.

Information that has been redacted is exempt under Section 40 (Personal Information) or Section 43 (Commercial Interests) of the Freedom of Information (FOI) Act and is therefore withheld.

Section 40 provides that personal information may be exempt from release where to do so would contravene data protection principles. Section 43 provides that information will be exempt from release where to do so would or would be likely to prejudice commercial interests. Furthermore, we do not believe that there is an overriding public interest in disclosing the redacted information in this instance.

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Yours sincerely,

FOI Team,
Vigilance and Risk Management of Medicines Division

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The Information Commissioner's Office
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Cheshire
SK9 5AF

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